

In the Claims:

Please amend the claims as follows:

1. (withdrawn) A device which, via at least one surface or one portion, is arranged to be applied to bone and/or tissue in the human body, for example jaw bone, and which, at the surface or portion, is provided with an agent which stimulates bone growth, preferably HA (hydroxyapatite), where at least one surface-bearing part or the portion comprises or consists of compressed bone-compatible and/or tissue-compatible material, preferably titanium powder, characterized in that the powder material and the bone-growth-stimulating agent form a composite material which is obtained by means of impact compaction and, if appropriate, sintering.

2. (withdrawn) The device as claimed in patent claim 1, characterized in that the bone-growth-stimulating/HA agent is arranged completely or partially in or at the actual surface layer and can thus be exposed to the bone and/or tissue in question.

3. (withdrawn) The device as claimed in patent claim 1, characterized in that the bone-growth-stimulating agent is in the form of particulate fractions with sizes in the range of 90-120  $\mu\text{m}$ .

4. (withdrawn) The device as claimed in patent claim 1, characterized in that titanium powder with considerable purity, preferably a purity of 99.99%, and a relatively small particle

size (Wah Chang HP (or CP) -325 Mesh T080014 (010607)) constitutes the base for the composite structure.

5. (withdrawn) The device as claimed in claim 1, characterized in that titanium powder in a quantity of ca. 90-98%, preferably ca. 95%, and HA powder in a quantity of 2-10%, preferably 5%, form the starting material for the material compacted by impactation and possible sintering.

6. (currently amended) A method for producing a device, ~~for example an implant,~~ which, via at least one surface or one portion, is arranged to be applied to bone and/or tissue in ~~the a~~ human body, ~~for example jaw bone, and which, the device~~ at the surface or portion, ~~is provided with an~~ comprising a powder agent which stimulates bone growth, ~~preferably HA,~~ ~~where wherein~~ at least one surface-bearing part or the portion ~~is made of~~ comprises compressed bone-compatible and/or tissue-compatible powder material, ~~preferably titanium powder,~~ the method comprising:

a) mixing together the bone-compatible and/or tissue-compatible powder material and said powder agent ~~which is in powder form,~~

b) applying the mixture in a mold cavity ~~belonging to of~~ a mold ~~applied~~ arranged in a machine which effects impact compaction ~~and which operates with a high impact compaction energy,~~

e) activating ~~the an~~ an impacting unit of the machine so that ~~it the impacting unit~~ the impacting unit acts on the mold and transfers the energy to the powder mixture and thereby creates a blank for the device, and

d) treating the blank in one or more treatment units for producing the device from the blank.

7. (currently amended) The method ~~as claimed in patent~~ according to claim 6, ~~characterized in that wherein treating the blank is comprises sintering sintered and/or heat-treated heat treating and is subjected~~ subjecting the blank to chemical, electrochemical and/or mechanical treatment or machining (~~milling, turning, shot-peening, etc.~~).

8. (currently amended) The method ~~as claimed in patent~~ according to claim 6, ~~characterized in that, in step a), wherein the bone-compatible and/or tissue-compatible powder material comprises~~ titanium powder ~~of considerable purity, for example 99.99%, and relatively small particle size is mixed together with HA, for example sintered HA, which and the powder agent comprises hydroxyapatite that~~ has been crushed and screened to ~~the~~ a fraction 90-120  $\mu\text{m}$ .

9. (currently amended) The method ~~as claimed in patent~~ according to claim 8, ~~characterized in that wherein the mixture consists of ea. comprises about 95% titanium powder and 5% HA hydroxyapatite powder, and wherein the titanium powder and the hydroxyapatite powder the powders are mixed in the~~ a dry state, with agitation and stirring.

10. (currently amended) The method ~~as claimed in patent~~ according to claim 8, ~~characterized in that wherein the machine is controlled so as to generate an impact compaction energy of ea. about 335 Nm or higher and to execute one or more impacts against the mold.~~

11. (currently amended) The method ~~as claimed in~~ according to claim 6, characterized ~~in that wherein particles of the titanium powder particles~~ are compressed to a substantial density, for example 98%, and ~~in that there is substantial surrounding of the HA~~ such that the particles of titanium powder substantially surround particles of the powder agent.

12. (currently amended) The method ~~as claimed in~~ according to claim 6, characterized ~~in that the wherein~~ positions of the HA particles of the powder agent ~~in the composite material~~ are controlled upon mixture and application in the mold cavity of the mold, and ~~in that wherein~~ the blank is machined so that HA particles of the powder agent are present at the surface exposed to the bone and/or tissue.

13. (cancelled)

14. (new) The method according to claim 6, wherein the powder agent comprises hydroxyapatite.

15. (new) The method according to claim 6, wherein the device comprises an implant.

16. (new) The method according to claim 6, wherein the device is applied to a jaw bone.

17. (new) The method according to claim 6, wherein the compressed bone-compatible and/or tissue-compatible powder material comprises titanium powder.

18. (new) The method according to claim 6, wherein the machining comprises milling, turning, or shot-peening.

19. (new) The method according to claim 6, wherein the titanium powder has a purity of 99.99%.